

OBJECTIVES: 1) To evaluate the relationship between asthma education and five measures of HRQoL—general health (GH), physical distress (PD), mental distress (MD), activity limitation (AL) and unhealthy days (UD), and 2) To find factors associated with receiving asthma education. **METHODS:** Data of adult US asthmatic individuals were obtained from the 2009 BRFSS ACBS. Asthma education comprises of five components: receiving asthma management plan, monitor peak flow, recognize early signs or symptoms of an asthma episodes, respond to early signs or symptoms of an episode, and change environment to improve asthma. Those who received all five components were labeled as “asthma education” group and those who received none of the five components were labeled as “no asthma education” group. Group comparisons for HRQoL measures were done by Chi-square test using SAS 9.2. **RESULTS:** From a total of 1472 individuals identified in this study, 54.55% did not receive all five components of asthma education. Individuals in the asthma education group reported significantly better GH (57.70% vs. 49.81%, $p=0.002$), fewer PD (64.28% vs. 59.15%, $p=0.046$) and fewer UD (52.32% vs. 44.46%, $p=0.002$). No significant differences were found for MD and AL due to high proportion of study individuals (>70%) reporting better HRQoL for these measures. Proportion of asthma education for all five components was significantly lower in; elderly, males, less than high school education, retired, and those with low income. **CONCLUSIONS:** Asthma education showed a positive impact on HRQoL measures – GH, PD and UD. Factors associated with non-receipt of all five asthma education components were age, gender, education, employment and income level. Interventions to encourage asthma education and overcome the inequality for the receipt of asthma education should be designed.

PHS67

HEALTH-RELATED QUALITY OF LIFE IN DIABETES

Kishore G¹, Nagaraj P², Ravouru N³¹Visveswarapura Institute of Pharmaceutical Sciences, Bangalore, India, ²Visveswarapura Institute of Pharmaceutical Sciences, Bangalore, India, ³SPMVV, Tirupati, India

OBJECTIVES: To study the health-related quality of life (HRQOL) of 300 patients with type 2 diabetes mellitus at a clinical setting of a tertiary care hospital. **METHODS:** The socio-demographic and clinical parameters of this group of patients were collected and their HRQOL was assessed using a generic questionnaire, the WHOQOL-BREF. **RESULTS:** The results revealed that in our study the patients were predominantly male (58%). The mean age of the study participants was found to be 56.97±11.57 years with 67.6% of the patients above the age of 50 years. Seventy-six percent of patients reported suffering from at least one other chronic non-diabetic medical condition like hypertension (63.33%) and/or hyperlipidemia (40.63%). Diabetic complications were also prevalent (62.4%). The highest mean domain score in our study population was in the social relationship domain (13.10±2.33), and the lowest was in the environment domain. The respondents with a lower family income, less education, increased age, female gender, single marital status and a rural background reported significantly lower HRQOL scores. **CONCLUSIONS:** Diabetes mellitus is a chronic disease of life long duration affecting about 9% of India's population. The number of people with diabetes is steadily increasing in India due to population growth, aging, urbanization and the increasing prevalence of obesity as well as physical inactivity. By identifying specific subsets of patients most likely to benefit from specialized care from health care providers, this study contributes by providing important data that may have direct implications for health care providers in their practice.

HEALTH SERVICES – Health Care Use & Policy Studies

PHS68

THE ROLE OF SOCIO-ECONOMIC FACTORS IN THE NON RENEWAL DECISION MAKING OF NATIONAL HEALTH INSURANCE POLICY HOLDERS IN THE BEKWAI MUNICIPAL

Agyei-Baffour P¹, Dankwa E²¹Kwame Nkrumah University of Science and Technology, Kumasi, Ghana, ²Ghana Health Service, Kokofu, Ghana

OBJECTIVES: To identify the socio-economic factors that influence the non renewal decision of National Health Insurance Policy Holders. **METHODS:** An analytical cross sectional survey was conducted. Four hundred and fifty-four (454) subjects from the non renewal list at the Bekwai Mutual Health Insurance Scheme were randomly selected and interviewed using validated structured and pre-tested questionnaires. A hand delivered questionnaire was used to collect data with the help of three trained research assistants after consenting process had been completed. The data was entered analyzed using SPSS version 16 and analysis was done for the various study variables. All statistical tests were run at 95% confidence interval and at 5% significance level. **RESULTS:** Two hundred two (44.5%) of the respondents reside in the urban setting while 252 (55.5%) of respondents was from the rural area. Eighty-seven out of the 454 respondents representing 19.2% were traders while 84 (18.5%), 80 (17.6%), 54 (11.9%) and 16 (3.5%) respectively. Most respondents were married suggesting that decoupling the exemption of children under 18 years old from parents has served as a disincentive to the registration of parents. Urban respondents who had no formal education and primary school level were 3.15 times and 2.32 times more likely not to renew their health insurance, as compared with rural respondents. Artisans, clericals and students were 0.80 times, 0.95 times, and 0.79 times less likely not to renew their health insurance respectively. Respondents who fall within income brackets 1-100, 101-200 and 201-300 were 0.54 times, 0.72 times and 0.78 times less likely not to renew health insurance. **CONCLUSIONS:** Socio-economic factors, such as education, income, marital status and employment had influence on the retention decisions. Implementing pro-poor social interventions and conducting further studies to access the influence of beliefs,

knowledge and status of health insurance on health care utilization could be helpful.

PHS69

COMPARISON BETWEEN THE UNITED STATES PREVENTIVE SERVICES TASK FORCE RECOMMENDATIONS AND MEDICARE COVERAGE POLICIES FOR PREVENTIVE SERVICES IN THE UNITED STATES

Garg V¹, Arabyat R¹, Raisch DW²¹University of New Mexico, Albuquerque, NM, USA, ²University of New Mexico College of Pharmacy, Albuquerque, NM, USA

OBJECTIVES: The Secretary of the Department of Health and Human Services is empowered by law to base Medicare reimbursement on the United States Preventive Services Task Force (USPSTF) recommendations (Grade A, benefits certainty high; B, benefits certainty moderate; and D, recommends against service). The purpose of this study was to evaluate the congruence between the USPSTF recommendations and their respective Medicare coverage policies. **METHODS:** The USPSTF website and Medicare preventive services guide were reviewed through December 2012 to study alignment between the USPSTF recommendations and Medicare coverage policies. Two reviewers independently extracted the data. We excluded USPSTF recommendations that were: I or C grade, inactive, for injury/violence, only valid for people <65 years, for preventive medication, or update-in-progress. Included USPSTF recommendations were categorized by therapeutic area (e.g., cancer), sub-therapeutic area (e.g., breast cancer), and recommendation class (i.e., screening and counselling). More than one recommendation for same sub-therapeutic area was treated separately. **RESULTS:** A total of 128 USPSTF recommendations were found, of which 37 (28.90%) were included in the analysis. The largest percentage of recommendations were for cancer (29.72%, n=11), followed by infectious diseases (27.02%, n=10). The vast majority were for screening (91.89%, n=34). Most of the Grade A and B USPSTF recommendations (94.11%, n=16) were covered by Medicare. However among USPSTF grade D recommendations, 35% (n=7) were also covered by Medicare (e.g., prostate-specific antigen based prostate cancer screening). As per the health care reform's objective of investment in prevention and illness, all of the included USPSTF recommendations were covered by Medicare without out-of-pocket payment, except one (i.e., tobacco cessation counselling). **CONCLUSIONS:** Medicare coverage policies for preventive services should be re-examined, especially those for which the USPSTF has issued grade D recommendations. Continuing such grade D recommendations without any established benefits will likely lead to unnecessary health care services and costs.

PHS70

KNOWLEDGE AND PERCEPTIONS OF PRESCRIBERS REGARDING ADHERENCE TO STANDARD TREATMENT GUIDELINES FOR MALARIA. A THREAT TO RATIONAL TREATMENT PRACTICES FOR MALARIA IN PAKISTAN

Malik M¹, Hassali MA², Shaffie A³, Hussain A⁴¹University Sains Malaysia, Hamdard University, Islamabad, Pakistan, ²Universiti Sains Malaysia, Penang, Malaysia, ³USM, Penang, Malaysia, ⁴Hamdard University, Islamabad, Pakistan

OBJECTIVES: To explore the perceptions and knowledge of prescribers towards adherence to standard treatment guidelines for malaria in two cities of Pakistan; Islamabad (national capital) and Rawalpindi (twin city). **METHODS:** A descriptive, cross sectional study design was used to evaluate the knowledge and perceptions of prescribers regarding adherence to standard treatment regimen for malaria. A questionnaire was developed through focused group discussions and distributed randomly to a sample of 360 prescribers in the two cities. Kruskal-Wallis test ($p\leq 0.05$) was used to compare the knowledge of prescribers (regarding standard malaria regimen) having different designations, level of experiences and their work affiliation with different levels of health care facilities. While Mann Whitney test ($p\leq 0.05$) was used to compare the knowledge among different genders of prescribers practicing in public and private health care facilities in the two cities. **RESULTS:** A significant difference ($p\leq 0.05$) among the knowledge of prescribers (regarding standard treatment regimen for malaria) having different designations, level of experience, working in different sectors and levels of health care facilities in the two cities was observed. However, no significant difference ($p\leq 0.05$) in the knowledge of different genders of prescribers was observed. **CONCLUSIONS:** The results of the present study showed that the overall knowledge of prescribers regarding standard treatment regimen for malaria in the two cities of Pakistan was inadequate. Most of the prescribers were unaware of correct standard treatment regimen for the treatment of Plasmodium falciparum and Plasmodium vivax.

PHS71

INTEGRATED CARE PROGRAMS IN EUROPE: FACTORS OF SUCCESS

Hoepfer K¹, Amelung VE¹, Hartmann J¹, Hermanowski T², Krauth C¹¹Hannover Medical School, Hannover, Germany, ²Department of Pharmacoeconomics, Medical University of Warsaw, Warsaw, Poland

OBJECTIVES: To determine the clinical and economic effects and components of integrated care programs to improve management of chronic diseases in Europe on the basis of systematic reviews and meta-analyses. **METHODS:** A systematic literature review was performed. Medline, Scopus, EMBASE, WHOLIS and The Cochrane Library of Systematic Reviews for English and German language articles were searched for English- and German language articles from 2002 to 2012. Articles were included if an explicit research question was defined, focussed on care that involves more than one provider, reported clinical outcomes of care, and employed acceptable experimental or quasi-experimental study designs as defined by the Cochrane Effective Practice and Organization of Care Group. More than 50% of included primary studies had to be RCT's and

conducted in Europe. **RESULTS:** The search identified 25,135 articles, 889 titles were included for further screening and 409 abstracts for the article review step. A total of 308 abstracts did not meet the inclusion/exclusion criteria, leaving a pool of 101 articles for full text evaluation. Overall, 10 systematic reviews were identified, with 238 primary studies, 129 conducted in Europe. The sample sizes from the included primary studies from Europe revealed 20 to 15,343 patients with a mean of 514. Chronic conditions investigated were: heart failure, chronic obstructive pulmonary disease, diabetes, and cancer. Of the outcomes more frequently studied, integrated care appeared to improve quality of life and reduce hospitalization. But often results remained inconclusive. **CONCLUSIONS:** Providing a conclusion across the different chronic conditions is not possible. Therefore, only disease specific conclusions can be drawn. Our review suggests that integrated care might be advantageous for specified groups of patients, e.g. heart failure. Furthermore, it remains unclear which specific component is associated with the highest benefit for patients across chronic conditions.

PHS72

AGE-RELATED EMERGENCY DEPARTMENT RELIANCE (EDR) AND HEALTH CARE RESOURCE UTILIZATION IN PATIENTS WITH SICKLE CELL DISEASE (SCD)

Blinder M¹, Vekeman F², Sasane M³, Trahey A⁴, Paley C³, Magestro M³, Duh MS⁴

¹Washington University in St. Louis, St. Louis, MO, USA, ²Groupe d'analyse, Montréal, QC, Canada, ³Novartis Pharmaceuticals Corporation, East Hanover, NJ, USA, ⁴Analysis Group, Inc., Boston, MA, USA

OBJECTIVES: For SCD patients, inadequate care during pediatric to adult transition may result in increased emergency department (ED) utilization. Emergency department reliance (EDR: total ED visits/total ambulatory [outpatient+ED] visits) identifies the proportion of ED visits in relation to all ambulatory visits. This study aimed at investigating age-related patterns of EDR and associated health care costs in SCD patients. **METHODS:** State Medicaid data from Florida, New Jersey, Missouri, Iowa, and Kansas were analyzed. Patients with ≥ 2 SCD diagnoses (ICD-9 282.6x) and ≥ 1 blood transfusion were included. Quarterly rates of EDR and SCD complication-related ED visits as well as health care costs were evaluated. Based on published thresholds, high EDR was defined as >0.33 . Regression analyses were used to assess risk factors for high EDR and calculate adjusted costs difference between patients with high versus low EDR. **RESULTS:** A total of 3208 patients were identified; mean (SD) observation period was 6.5 (3.2) years. Mean ED visits/quarter increased from 0.76 to 2.23 between age 15 and 23, reaching a peak of 2.9 at age 36. The most common SCD complication-related ED visits were pain, infection, and pneumonia. EDR rose from 0.15 to 0.29 between age 15 and 23, and remained high thereafter. Patients were more likely to have high EDR during the post-transition period (≥ 18 years old, odds ratio [OR]: 2.38, $p < 0.001$) and when experiencing an SCD complication (OR: 4.18, $p < 0.001$). Patients with high EDR incurred higher inpatient and ED costs, resulting in higher total costs (high vs. low EDR, adjusted costs difference, OP: -\$285; IP: \$3,485; ED: \$120; Rx: -\$91; total: \$3,086, $p < 0.001$ for all). **CONCLUSIONS:** Compared to children, SCD patients transitioning to adulthood relied more on ED for their care and those with high EDR incurred higher health care costs, highlighting the need to improve access to care for transitioning and adult SCD patients.

PHS73

PHARMACIST-LED SERVICES TO PATIENTS WITH RESPIRATORY DISEASES: FEASIBLE FROM A QUALITY AND REIMBURSEMENT PERSPECTIVE?

Willey VI¹, Simon S¹, Akkineni S¹, Reinhold JA¹, Kelly BL², Kim EA², Willey KH², Cawley MJ¹

¹University of the Sciences, Philadelphia, PA, USA, ²Quality Family Physicians, Hockessin, DE, USA

OBJECTIVES: Pharmacists are qualified to provide many services that are core to integrated care models. Expanding services to diverse patient populations will increase pharmacists' value. This study describes the experience, preliminary outcomes and revenue model justification associated with the implementation of pharmacist-led care for patients with respiratory disorders. **METHODS:** Medical and billing record review was performed on patients with respiratory symptoms referred to the pharmacist from May 2011 to September 2012 within a community-based, medical home, primary care practice. Patients referred were those with respiratory symptoms in which the physician sought objective lung function data and additional support to assist in properly diagnosing and treating the patient. Pharmacist interventions included collection of a detailed pulmonary and medication history, spirometry, and on applicable patients, disease state education, medication care plans and device education, and smoking cessation. Outcomes described included quality and results of spirometry testing, pharmacist recommendations, recommendations for specialist care and payment for services. **RESULTS:** Thirty-four patients (76.5% female; mean age=49.6 \pm 17.6) were seen by the pharmacist and assessed by spirometry. Spirometry met American Thoracic Society quality measures in 82.5% of tests with the following results: 64.7% normal, obstruction (8.8% mild, 14.7% moderate, 2.9% severe), and 8.8% probable restriction. Pharmacist recommendations that were implemented included the use of short-acting-beta-agonists (23.5%), corticosteroids (20.6%), anti-cholinergics (14.7%), and long-acting-beta-agonists (11.8%). Smoking cessation was recommended for 11.8% of patients and 44.1% received specialist referrals. The mean overall payment for the services provided at these visits was \$144.43 \pm 36.34. **CONCLUSIONS:** Pharmacist involvement in the care of patients with respiratory disorders provided valuable, quality lung function data and care plan recommendations to physicians and education to patients. These preliminary results support the pharmacist expanding their role in the medical home by providing physician/patient care services, including spirometry, to patients with respiratory disorders from both a clinical and economic perspective.

PHS74

DRUG-RELATED PROBLEMS AND MEDICATION ERRORS: A LITERATURE REVIEW ON ECONOMIC OUTCOMES IN SUB-SAHARAN AFRICA

McRae J¹, Lovett A¹, Ohaya V¹, Ohaya J¹, Pounds T²

¹Mercer University, Atlanta, GA, USA, ²Atlanta Medical Center, Atlanta, GA, USA

OBJECTIVES: To review the literature published within the last decade related to drug-related problems and medication errors in Sub-Saharan Africa. This article provides a discussion on pharmaceutical care, with a focus on economic outcomes. **METHODS:** A search using Medline, Embase was conducted over the timeframe January 2002–December 2012 using key words such as: Sub-Saharan Africa, medication errors, economics, and pharmaceutical care. The abstract and/or full text of each article was reviewed. **RESULTS:** Twenty studies were identified for review. The most common problems in the pharmacy system were improper labeling, counterfeit drugs, lack of patient education, and inadequate drug distribution. The lack of electronic medical records and payment systems prevent the assessment of clinical cost outcomes. In a study conducted by the University of Benin, of 1500 pharmacists, 93% reported that they would be willing to participate in "any training program to enable them to practice pharmaceutical care." There is a lack of pharmacists able to provide pharmaceutical care as defined as the direct, responsible provision of medication-related care designed to achieve definite outcomes. The shortage of pharmacists is due to few training institutions, migration, inadequate pay and poor working conditions. Specifically, 25% of the world's global burden of disease is in Sub-Saharan Africa, while this area comprises 3% of the world's health workers. These factors contribute to an increase in medication errors. **CONCLUSIONS:** Sub-Saharan Africa lacks the necessary governmental regulation to ensure a decrease in medication errors. The government may consider streamlining their drug distribution system through the enforcement of regulations and the use of information technology in the health care delivery system. Additional studies are needed to examine economic outcomes. Several studies provide information on cost effectiveness and quality of life, but these studies are specific to the HIV/AIDS population.

PHS75

HEALTH CARE FRAUD 2006 TO 2011

Lu K¹, Chen B², Qureshi Z³, Xirasagar S⁴, Sartor O⁴, Bennett C²

¹South Carolina College of Pharmacy – USC Campus, Columbia, SC, USA, ²University of South Carolina, Columbia, SC, USA, ³Arnold School of Public Health, University of South Carolina, Columbia, SC, USA, ⁴Tulane University, New Orleans, LA, USA

OBJECTIVES: Health care fraud is a long-standing problem, accounting for \$75 billion in 2009. Congress amended the False Claims Act (FCA) in 1986 to allow qui tam relators ("whistleblowers") to receive up to 30% of anti-fraud recoveries. Most studies investigate health care fraud involving the pharmaceutical industry, so it has not been possible to contextualize fraud involving health care sectors other than the pharmaceutical industry. Herein, we review all recently concluded major federal health care fraud investigations. **METHODS:** All cases involved health care corporations and federal FCA. Data were collected from Lexis/Nexis News (search terms: "Health care fraud", "False Claims Act" and "Qui tam"), the Taxpayers against Fraud and the DOJ websites (2006-2011). Only cases with recoveries over \$5 million ("major cases") were included. Data were abstracted on allegations, financial settlements, occupations of and payments to qui tam relators. Cases are reported separately as qui tam- versus non-qui tam-initiated to document whether the Congressional intent to encourage whistleblowing achieved its intended goal. **RESULTS:** Between 2006 and 2011, 123 major qui tam health care FCA cases concluded, totaling \$15.7 billion in recoveries (mean recovery: \$128 million). Billing fraud, kickbacks, off-label marketing, and marketing unsafe pharmaceuticals were the most commonly implicated activities. Pharmaceutical manufacturers accounted for 31% of, and \$11.3 billion (70%) in recoveries among qui tam relator cases. Also, 52 non-qui tam cases closed in this 5-year period, totaling \$3.7 billion in recoveries (mean recovery: \$71 million). Implicated activities included fraudulent billing, inappropriate financial relationships, off-label marketing, or marketing unsafe pharmaceuticals. **CONCLUSIONS:** In conclusion, federal investigations of fraud and abuse involving health care are increasing in both depth and breadth, and qui tam relators have an important role in detecting important fraud and abuse.

PHS76

ANALYSIS OF 2011 MEDICAID FEE-FOR-SERVICE OUTPATIENT DRUG UTILIZATION, EXPENDITURES AND PHARMACY REIMBURSEMENT RATES

Alshehri A¹, Balkhi B¹, Seoane-Vazquez E¹, Szeinbach SL²

¹International Center for Pharmaceutical Economics and Policy, Massachusetts College of Pharmacy and Health Sciences, Boston, MA, USA, ²Ohio State University, Columbus, OH, USA

OBJECTIVES: The Patient Protection and Affordable Care Act will expand the eligibility of the Medicaid program to millions of Americans in 2014. Utilizing more generic drugs and setting appropriate pharmacy reimbursement rates could result in substantial savings to the Medicaid program. This study assessed 2011 state-level, fee-for-service Medicaid generic and brand drug utilization and expenditures, and pharmacy reimbursement rates. **METHODS:** Medicaid fee-for-service outpatient pharmacy utilization and expenditures, and reimbursement rates (ingredient cost and dispensing fees) for the year 2011 were extracted from state-level data provided from the Centers for Medicare and Medicaid Services. Descriptive analyses were performed for all variables in the data set. Linear regression analysis was performed to assess the relationship between ingredient cost, dispensing fees and drug utilization. The significance level for variables was 0.05. **RESULTS:** Fee-for-service Medicaid expenditures (n=46 states) reached \$27.8 billion with drug utilization accounting for 173.4 million claims in 2011. Generic expenditures represented 17.3% of total expenditures (range=10.3%-29.2%) and